Docket No. 1377-0170P

## IN THE CLAIMS

1. (currently amended) A method for identifying a gene having a role in which may be involved with the presentation of diabetic nephropathy, which method comprises

culturing mesangial cells in a medium in the presence of transforming growth factor £1 (TGF-£1) and a concentration of glucose sufficient to induce differential expression of gene susceptible to such differential expression;

and identifying the gene so induced by suppression subtractive hybridization.

- 2. (original) A method according to Claim 1, wherein the mesangial cells are cultured in the presence of a concentration of glucose sufficient to induce up-regulation of a gene susceptible to such up-regulation.
- 3. (previously amended) A method according to Claim 1, wherein the concentration of glucose is greater than 5 mM.
- 4. (previously amended) A method according to Claim 1, wherein the mesangial cells are subjected to mechanical strain.
  - 5. (cancelled)

Docket No. 1377-0170P

- 6. (previously amended) A method according to Claim 1, wherein the possibility of differential expression due to hyperosmolarity is excluded.
- 7. (currently amended) A method according to Claim 1, wherein the gene so differently expressed is a gene of SEQ ID NO:1 which includes a sequence selected from the group consisting of:
  - 1) SEQ ID NOS:-1-3;
  - 2) SEQ ID NO:4;
  - 3) SEQ ID No:5; and
  - 4) SEQ ID NO:6.
- 8. (previously amended, withdrawn) Use of a gene identified by a method according to Claim 1, as a diagnostic marker for the progression and presentation of diabetic nephropathy.
- 9. (previously amended, withdrawn) Use of a gene identified by a method according to Claim 1, as an index of disease activity and the rate of progression of diabetic nephropathy.
- 10. (previously amended, withdrawn) Use of a gene identified by a method according to Claim 1, as a basis for identifying drugs for use in the prevention and/or therapy of diabetic nephropathy.

Docket No. 1377-0170P

11. (currently amended) A An isolated gene encoded by a sequence of SEQ ID NO:1 selected from any one of SEQ IDS NOS:1 3, 5 and 6 according to Claim 7.